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OM protein - protein search, using sw model

Run on: November 5, 2004, 13:50:17 ; Search time 74.3927 seconds
(without alignments)
3081.324 Million cell updates/sec

Title: US-09-937-636-4

Perfect score: 3377
Sequence: 1 MLLPILLSLLGSGQAMGR.....RPEARNPKGTADYAEVKFQ 639

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 2002273 seqs, 358729299 residues

Total number of hits satisfying chosen parameters: 2002273

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :
1: Genesecp23Sep04:*
2: Genesecp1980s:*
3: Genesecp1980s:*
4: Genesecp2000s:*
5: Genesecp2001s:*
6: Genesecp2002s:*
7: Genesecp2003as:*
8: Genesecp2003bs:*
9: Genesecp2004s:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	3377	100.0	639	2	AAW81023 Human sia
2	3377	100.0	639	3	AAY97543 Human obe
3	3354	99.3	639	3	AAB25580 CD33-like
4	3354	99.3	639	6	ADA27052 Human nov
5	3354	99.3	639	8	AD86582 Novel hum
6	3338	98.8	637	5	AAU87087 Sialic ac
7	3334	98.7	637	5	ADA27153 Human nov
8	3334	98.7	637	7	ADD26582 Siglec-10
9	3334	98.7	637	8	ADI37010 Novel hum
10	3334	98.7	637	5	ADL82805 Human PRO
11	3299	97.7	627	5	AAU87075 Sialic ac
12	3174.5	94.0	710	7	ADD19314 Human sec
13	2903.5	83.0	544	5	AAU87074 Sialic ac
14	2900.5	82.6	544	2	AAU87074 Human PRO
15	2900.5	82.6	544	3	AAU87074 Human PRO
16	2900.5	82.6	544	6	AAU87074 Human PRO
17	2900.5	82.6	544	6	AAU87074 Human PRO
18	2900.5	82.6	544	6	AAU87074 Human PRO
19	2900.5	82.6	544	6	AAU87074 Human PRO
20	2900.5	82.6	544	6	AAU87074 Human PRO
21	2900.5	82.6	544	6	AAU87074 Human PRO
22	2900.5	82.6	544	6	AAU87074 Human PRO
23	2900.5	82.6	544	6	AAU87074 Human PRO
24	2900.5	82.6	544	6	AAU87074 Human PRO
25	2900.5	82.6	544	6	AAU87074 Human PRO

26	2790.5	82.6	544	6	ABU96187	Novel hum
27	2790.5	82.6	544	6	ABU92618	Human sec
28	2790.5	82.6	544	6	ABO08695	Human sec
29	2790.5	82.6	544	6	ABO02747	Human sec
30	2790.5	82.6	544	6	ABR74901	Human sec
31	2790.5	82.6	544	6	ABR94663	Human sec
32	2790.5	82.6	544	6	ABO25226	Novel hum
33	2790.5	82.6	544	6	ABU85636	Human PRO
34	2790.5	82.6	544	6	ABU98796	Novel hum
35	2790.5	82.6	544	6	ABU98011	Novel hum
36	2790.5	82.6	544	6	ABU91717	Novel hum
37	2790.5	82.6	544	6	ABU72232	Novel hum
38	2790.5	82.6	544	6	ABU89410	Human PRO
39	2790.5	82.6	544	6	ABU86251	Human sec
40	2790.5	82.6	544	6	ABU67464	Human sec
41	2790.5	82.6	544	6	ABU80492	Human PRO
42	2790.5	82.6	544	6	ABR99410	Human sec
43	2790.5	82.6	544	6	ABR98800	Human sec
44	2790.5	82.6	544	6	ABO16323	Human sec
45	2790.5	82.6	544	6	ABR92223	Human sec

ALIGNMENTS

RESULT 1
AAW81023
ID AAW81023 standard; protein; 639 AA.
XX
AC AAW81023;
XX
DT 26-APR-1999 (first entry)
XX
DE Human sialoadhesin family 4 (SAF-1) polypeptide.

SAF-4; sialoadhesin family; human; therapy; diagnosis; cancer;
inflammation; autoimmune disease; allergy; asthma; inflammation;
cerebellar degeneration; Alzheimer's disease; Parkinson's disease;
multiple sclerosis; amyotrophic lateral sclerosis; head injury;
septic shock; sepsis; stroke; osteoporosis; osteoarthritis;
ischemia reperfusion injury; cardiovascular disease; kidney disease;
liver disease; myocardial infarction; hypertension; hypotension; AIDS;
myelodysplastic syndrome; aplastic anaemia; baldness; infection.

XX Homo sapiens.

XX WO9853840-A1.

XX 03-DEC-1998.

XX 27-MAY-1998; 98WO-US010791.

XX 27-MAY-1997; 97US-0047572P.

XX (SMIK) SMITHKLINE BEECHAM CORP.

XX Kikly KK, Erickson-Miller CL;

XX WPI; 1999-080779/07.

XX N-PSDB; AAV99911.

XX New sialoadhesin family 4 polypeptides and polynucleotides - useful to treat various diseases associated with SAF-4 expression.

XX Claim 1; Page 31; 48pp; English.

XX This is the amino acid sequence of new human sialoadhesin family 4 (SAF-4), as deduced from the nucleotide sequence of an isolated cDNA clone (see AAV99911). SAF-4 polynucleotides and polypeptides, and methods for producing such polypeptides in transformed host cells using recombinant techniques are disclosed. SAF-4, its agonists and antagonists, and nucleic acid molecules that enhance or inhibit SAF-4 expression, may be used to treat patients in need of enhancement or inhibition of SAF-4

expression or activity. Conditions that may benefit from such treatment include cancer, inflammation, autoimmunity, allergy, asthma, rheumatoid arthritis, CNS inflammation, cerebellar degeneration, Alzheimer's disease, Parkinson's disease, multiple sclerosis, amyotrophic lateral sclerosis, head injury damage and other neurological disorders, septic shock, sepsis, stroke, osteoporosis, osteoarthritis, ischemia reperfusion injury, cardiovascular disease, kidney disease, liver disease, ischemic injury, myocardial infarction, hypotension, hypertension, AIDS, myelodysplastic syndromes and other haematologic abnormalities, aplastic anaemia, male baldness pattern and bacterial, protozoal, fungal and viral infections related to SAP-4 polypeptide activity. Methods of identifying agonists, antagonists/inhibitors are also provided, as well as diagnostic assays for detecting diseases associated with inappropriate SAP-4 activity or levels

SQ Sequence 639 AA;

Query Match 100.0%; Score 3377; DB 2; Length 639;
Best Local Similarity 100.0%; Pred. No. 2.3e-232; Indels 0; Gaps 0;
Matches 639; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 MLLPILLSLLGGSGQMDGRFWIRVQESVMVPEGLCISVPCSPSYPRQDMTGSTPAYGYW 60
Db 1 MLLPILLSLLGGSGQMDGRFWIRVQESVMVPEGLCISVPCSPSYPRQDMTGSTPAYGYW 60
Qy 61 FKAVTTTKGAPVATNHQSRVEMSTRGEFOLTGDPKAGNSLVIRDAQMDQESQYFFRV 120
Db 61 FKAVTTTKGAPVATNHQSRVEMSTRGEFOLTGDPKAGNSLVIRDAQMDQESQYFFRV 120
Qy 121 ERGSVRYNFMNDGFFLKVTLSFTPRPDHNTDLTCHVDSRKGVSQAQRTVRLVAYAP 180
Db 121 ERGSVRYNFMNDGFFLKVTLSFTPRPDHNTDLTCHVDSRKGVSQAQRTVRLVAYAP 180
Qy 181 RDLVTSISRDNTPALEPQGNVPLEAKQGFLELLCAADSPATLSWLNQVLS 240
Db 181 RDLVTSISRDNTPALEPQGNVPLEAKQGFLELLCAADSPATLSWLNQVLS 240
Qy 241 HPWGPRLGLELPGVKAGDSGRYTCRAENRLGSGQRAALDLSVQYPPENLRVWVSQANRTV 300
Db 241 HPWGPRLGLELPGVKAGDSGRYTCRAENRLGSGQRAALDLSVQYPPENLRVWVSQANRTV 300
Qy 301 LENLGNCTSLPVLKQSLCIVCTHSSPPARLSWTQRCQVLSPSQSPDGVLELPRVQVE 360
Db 301 LENLGNCTSLPVLKQSLCIVCTHSSPPARLSWTQRCQVLSPSQSPDGVLELPRVQVE 360
Qy 361 HEGEFTCHARHPLGSGQVSLSVHYSPKLLGPGSCSWEAEGHLCSSQASPAFLSLANWL 420
Db 361 HEGEFTCHARHPLGSGQVSLSVHYSPKLLGPGSCSWEAEGHLCSSQASPAFLSLANWL 420
Qy 421 GEEILLEGNSQDSFEVTPSSAGPWANGSLSLHGGSLSGRLRCCEANVHGAQSGSILQLP 480
Db 421 GEEILLEGNSQDSFEVTPSSAGPWANGSLSLHGGSLSGRLRCCEANVHGAQSGSILQLP 480
Qy 481 DKKGLISTAFSNGAFPLGIGITALLFLCTALIMKILPKRPTOTETPRPSRHSHTILDYI 540
Db 481 DKKGLISTAFSNGAFPLGIGITALLFLCTALIMKILPKRPTOTETPRPSRHSHTILDYI 540
Qy 541 NVVPTAGLAQRNKAQTPNSPRTPLPGAPSPESKKNQKQYQLPSFPKSSTOAPES 600
Db 541 NVVPTAGLAQRNKAQTPNSPRTPLPGAPSPESKKNQKQYQLPSFPKSSTOAPES 600
Qy 601 QESCEELHYATLNPFGVYRPREARMPKGTQADYAEVKFO 639
Db 601 QESCEELHYATLNPFGVYRPREARMPKGTQADYAEVKFO 639

RESULT 2
ID AAY97543
XX AAY97543 standard; protein; 639 AA.
XX AC AAY97543;
XX 12-FEB-2001 (first entry)

XX Human obesity protein binding protein-2 homologue #2.
DB Human obesity protein binding protein-2 homologue; hOB-BP2h; obesity;
XX Obesity-related disorder; therapy.
XX Homo sapiens.
XX WO200059942-A2.
XX 12-OCT-2000.
XX 22-MAR-2000; 2000WO-US006682.
XX 02-APR-1999; 99US-0127667P.
XX (ELIL) LILLY & CO ELI.
XX Su EW, Wei J;
XX WPI; 2000-664992/64.
XX N-PSDB; AAA37848.
XX New human obesity protein binding protein-2 homologue nucleic acids,
XX polynucleotides and polypeptides useful for producing medicament for
XX treating obesity and/or obesity-related disorders.
XX Claim 9; Page 89-91; 92pp; English.
XX This sequence is a human obesity protein binding protein-2 homologue (hOB
XX -BP2h) of the invention. The hOB-BP2h nucleic acids and polypeptides may
XX be used for the manufacture of a medicament for the treatment of obesity
XX and/or obesity-related disorders. The hOB-BP2h nucleic acids are useful
XX as probes or amplification primers in the detection, quantification or
XX isolation of gene sequences or transcripts, for recombinant expression of
XX hOB-BP2h polypeptides, as immunogens in the preparation and screening of
XX antibodies, and in sense or antisense suppression of one or more hOB-BP2h
XX genes or nucleic acids, host cell or tissue in vivo or in vitro.
XX Antigenic epitope-bearing peptides and polypeptides are useful for
XX raising or screening antibodies that specifically binds to the hOB-BP2h
XX polypeptides

SQ Sequence 639 AA;

Query Match 100.0%; Score 3377; DB 3; Length 639;
Best Local Similarity 100.0%; Pred. No. 2.3e-232; Indels 0; Gaps 0;
Matches 639; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 MLLPILLSLLGGSGQMDGRFWIRVQESVMVPEGLCISVPCSPSYPRQDMTGSTPAYGYW 60
Db 1 MLLPILLSLLGGSGQMDGRFWIRVQESVMVPEGLCISVPCSPSYPRQDMTGSTPAYGYW 60
Qy 61 FKAVTTTKGAPVATNHQSRVEMSTRGEFOLTGDPKAGNSLVIRDAQMDQESQYFFRV 120
Db 61 FKAVTTTKGAPVATNHQSRVEMSTRGEFOLTGDPKAGNSLVIRDAQMDQESQYFFRV 120
Qy 121 ERGSVRYNFMNDGFFLKVTLSFTPRPDHNTDLTCHVDSRKGVSQAQRTVRLVAYAP 180
Db 121 ERGSVRYNFMNDGFFLKVTLSFTPRPDHNTDLTCHVDSRKGVSQAQRTVRLVAYAP 180
Qy 181 RDLVTSISRDNTPALEPQGNVPLEAKQGFLELLCAADSPATLSWLNQVLS 240
Db 181 RDLVTSISRDNTPALEPQGNVPLEAKQGFLELLCAADSPATLSWLNQVLS 240
Qy 241 HPWGPRLGLELPGVKAGDSGRYTCRAENRLGSGQRAALDLSVQYPPENLRVWVSQANRTV 300
Db 241 HPWGPRLGLELPGVKAGDSGRYTCRAENRLGSGQRAALDLSVQYPPENLRVWVSQANRTV 300
Qy 301 LENLGNCTSLPVLKQSLCIVCTHSSPPARLSWTQRCQVLSPSQSPDGVLELPRVQVE 360
Db 301 LENLGNCTSLPVLKQSLCIVCTHSSPPARLSWTQRCQVLSPSQSPDGVLELPRVQVE 360
Qy 361 HEGEFTCHARHPLGSGQVSLSVHYSPKLLGPGSCSWEAEGHLCSSQASPAFLSLANWL 420

AAV99912
ID AAV99912 standard; cDNA; 3099 BP.
XX
XX AC AAV99912;
XX
XX DT 26-APR-1999 (first entry)
XX
XX DS Human sialoadhesin family 4 (SAP-1) related EST clone.
XX
XX KW SAP-4; sialoadhesin family; human; therapy; diagnosis; cancer;
XX KW inflammation; autoimmune disease; allergy; asthma; inflammation;
XX KW cerebellar degeneration; Alzheimer's disease; parkinson's disease;
XX KW multiple sclerosis; amyotrophic lateral sclerosis; head injury;
XX KW septic shock; sepsis; stroke; osteoporosis; osteoarthritis;
XX KW ischemia reperfusion injury; cardiovascular disease; kidney disease;
XX KW liver disease; myocardial infarction; hypotension; hypertension; AIDS;
XX KW myelodysplastic syndrome; aplastic anaemia; baldness; infection; ss.
XX
XX OS Homo sapiens.
XX
XX FH Key Location/Qualifiers
XX CH 51.1249
XX FT /*tag a
XX FT /transl_except= (pos:1176..1177, aa:Pro)
XX FT /note= "this codon has an apparent 1 nucleotide deletion
XX FT which alters the reading frame"
XX
XX PN WO9853840-A1.
XX
XX PD 03-DEC-1998.
XX
XX PF 27-MAY-1998; 98WO-US010791.
XX
XX PR 27-MAY-1997; 97US-0047572P.
XX
XX PA (SMIX) SMITHKLINE BEECHAM CORP.
XX
XX PI Kikly KK, Erickson-Willer CL;
XX
XX DR WPI: 1999-080779/07.
XX
XX P-PSDB; AAW81024.
XX
XX PT New sialoadhesin family 4 polypeptides and polynucleotides - useful to
XX PT treat various diseases associated with SAP-4 expression.
XX
XX PS Claim 13; Page 31-32; 48pp; English.
XX
XX CC This is the nucleotide sequence of an expressed sequence tag (EST) clone
XX CC that encodes a polypeptide (see AAW81024) which shows close homology
XX CC and/or structural similarity (e.g. a conservative amino acid difference)
XX CC to new human sialoadhesin family 4 (SAP-4) polypeptide (see AAW81023).
XX CC SAP-4 polynucleotides and polypeptides, and methods for producing such
XX CC polypeptides in transformed host cells using recombinant techniques, are
XX CC disclosed. SAP-4, its agonists and antagonists, and nucleic acid
XX CC molecules that enhance or inhibit SAP-4 expression, may be used to treat
XX CC patients in need of enhancement or inhibition of SAP-4 expression or
XX CC activity. Conditions that may benefit from such treatment include cancer,
XX CC inflammation, autoimmunity, allergy, asthma, rheumatoid arthritis, CNS
XX CC disease, multiple sclerosis, amyotrophic lateral sclerosis, head injury
XX CC damage and other neurological disorders, septic shock, sepsis, stroke,
XX CC osteoporosis, osteoarthritis, ischemia reperfusion injury, cardiovascular
XX CC disease, kidney disease, liver disease, ischemic injury, myocardial
XX CC infarction, hypotension, hypertension, AIDS, myelodysplastic syndromes
XX CC and other haematologic abnormalities, aplastic anaemia, male baldness
XX CC pattern and bacterial, protozoal, fungal and viral infections related to
XX CC SAP-4 polypeptide activity. Methods of identifying agonists, and
XX CC antagonists/inhibitors are also provided as well as diagnostic assays
XX CC for detecting diseases associated with inappropriate SAP-4 activity or
XX CC levels
XX
XX 3Q Sequence 3099 BP; 770 A; 907 C; 813 G; 607 T; 0 U; 2 Other;

1071	DB		CTGAGCCCTTC	CAGGCGCTC	CAGACCCCGGGTCT	CTGAGCTGCCTCGGTTCAAGTGGAG	1130
1081	QY		CACGAAGAGAGTT	CACTCCCAAGCT	CTGGCAACCACTGGGCTCCAGCAAGTCTCTCTC	1140	
1131	DB		CACGAAGAGAGTT	CACTCCCAAGCT	CTGGCAACCACTGGGCTCCAGCAAGTCTCTCTC	1190	
1141	QY		AGCCTCTCCGTG	CACACTACTCCCGGAAGCT	GTGGGCGCTCTCTGCTCTCTGSGAGGCTGAG	1200	
1191	DB		AGCTCTCCGTG	CACACTACTCCCGGAAGCT	GTGGGCGCTCTCTGCTCTCTGSGAGGCTGAG	1250	
1201	QY		GGTGTGCACTG	CAGCTGCTCTCCAGGCGCAGCGCGGCCCTCTCTCTGCGCTGGTGGCTT	1260		
1251	DB		GGTGTGCACTG	CAGCTGCTCTCCAGGCGCAGCGCGGCCCTCTCTCTGCGCTGGTGGCTT	1310		
1261	QY		GGGAGGAGAGT	GTCTGGAAGGGAACAGCAGCCAGACTCTCTTCGAGGTCACCCCAAGTCTCA	1320		
1311	DB		GGGAGGAGAGT	GTCTGGAAGGGAACAGCAGCCAGACTCTCTTCGAGGTCACCCCAAGTCTCA	1370		
1321	QY		GCCGGCGCTTG	GGGCACACAGCTCCCTGAGCTTCCATGAGGCTCAGCTCCGGCTCAGG	1380		
1371	DB		GCCGGCGCTTG	GGGCACACAGCTCCCTGAGCTTCCATGAGGCTCAGCTCCGGCTCAGG	1430		
1381	QY		CTCCGCTGTG	AGCGCTGGAAAGTCCCATGAGGGGCCAGAGTGAATCCATCTGCAAGTCCCA	1440		
1431	DB		CTCCGCTGTG	AGCGCTGGAAAGTCCCATGAGGGGCCAGAGTGAATCCATCTGCAAGTCCCA	1490		
1441	QY		GATAAGAAGAG	ACTCATCTCAACGGCAATTTCTCAACGGAGCGTTTCTGGGAATCGGCATC	1500		
1491	DB		GATAAGAAGAG	ACTCATCTCAACGGCAATTTCTCAACGGAGCGTTTCTGGGAATCGGCATC	1550		
1501	QY		ACGGCTCTCTT	TTCTCTGCTGCTGGCGCTGATGATCTGAGATTTCTACGAGAGACGG	1560		
1551	DB		ACGGCTCTCTT	TTCTCTGCTGCTGGCGCTGATGATCTGAGATTTCTACGAGAGACGG	1610		
1561	QY		ACTCAGACAGAA	ACCCCGAGGCCAGGTTCTCCCGGCACAGCAGATCTCTGGATTATCATC	1620		
1611	DB		ACTCAGACAGAA	ACCCCGAGGCCAGGTTCTCCCGGCACAGCAGATCTCTGGATTATCATC	1670		
1621	QY		AAGTGTGTCCG	AGCGGTGCGCCCTGCGCTCAGAGCGGATCTCAGAAAGCCACACCAAC	1680		
1671	DB		AAGTGTGTCCG	AGCGGTGCGCCCTGCGCTCAGAGCGGATCTCAGAAAGCCACACCAAC	1730		
1681	QY		AGTCTCTCGG	ACCCCTTCCACAGGTGCTCTCCCGAGATCAAAAGAGAACACAGAAA	1740		
1731	DB		AGTCTCTCGG	ACCCCTTCCACAGGTGCTCTCCCGAGATCAAAAGAGAACACAGAAA	1790		
1741	QY		AAGCAGTATC	AGTGTGCCAGTCTTCCAGAACCCAAATCATCTCAGTCCAGGCCAGATCC	1800		
1791	DB		AAGCAGTATC	AGTGTGCCAGTCTTCCAGAACCCAAATCATCTCAGTCCAGGCCAGATCC	1850		
1801	QY		CAGAGAGACCA	AGAGAGTCCATTATGCAAGCTCAACTTCCAGGGGTCAAGCCAGG	1860		
1851	DB		CAGAGAGACCA	AGAGAGTCCATTATGCAAGCTCAACTTCCAGGGGTCAAGCCAGG	1910		
1861	QY		CCTGAGCCCGG	ATGCCCAGAGGCCACCCAGGGGGATTATGAGAGTCAAGTTCCAA	1917		
1911	DB		CCTGAGCCCGG	ATGCCCAGAGGCCACCCAGGGGGATTATGAGAGTCAAGTTCCAA	1967		

RESULT 3
AAV99911
ID AAV99911 standard; cDNA: 3099 BP.

AAV99911;

DT 26-APR-1999 (first entry)

Human sialoadhesin family 4 (SAF-1) cDNA.

SAF-4; sialoadhesin family; human; therapy; diagnosis; cancer; inflammation; autoimmune disease; allergy; asthma; inflammation; cerebellar degeneration; Alzheimer's disease; Parkinson's disease;

KW	multiple sclerosis, amyotrophic lateral sclerosis; head injury;
KW	septic shock; sepsis; stroke; osteoporosis; osteoarthritis;
KW	ischemia reperfusion injury; cardiovascular disease; kidney disease;
KW	liver disease; myocardial infarction; hypotension; hypertension; AIDS;
KW	myelodysplastic syndrome; aplastic anaemia; baldness; infection; ss.
XX	
XX	
XX	OS
XX	Homo sapiens.
XX	
XX	Key
XX	Location/Qualifiers
XX	51. 1970
XX	/*tag= a
XX	
XX	WO9853840-A1.
XX	
XX	03-DEC-1998.
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XX	27-MAY-1998; 98WO-US010791.
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XX	27-MAY-1997; 97US-0047572P.
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XX	(SMIX) SMITHKLINE BEECHAM CORP.
XX	
XX	Kirkly KK, Erickson-Miller CL;
XX	
XX	WPI; 1999-080779/07.
XX	
XX	9-ESDB; AAN91023.
XX	
XX	New sialoadhesin family 4 polypeptides and polynucleotides - useful to
XX	treat various diseases associated with SAR-4 expression.
XX	
XX	Claim 2; Page 30-31; 48pp; English
XX	

This cDNA clone codes for new human sialoadhesin family 4 (SAP-4) polypeptide (see AA081023). It can be obtained e.g. from a cDNA library derived from mRNA in cells of human primary dendritic cells, using expressed sequence tag analysis. SAP-4 polynucleotides and polypeptides, and methods for producing such polypeptides by recombinant techniques are disclosed. SAP-4, its agonists and antagonists, and nucleic acid molecules that enhance or inhibit SAP-4 expression, may be used to treat patients in need of enhancement or inhibition of SAP-4 expression or activity. Conditions that may benefit from such treatment include cancer, inflammation, autoimmunity, allergy, asthma, rheumatoid arthritis, CNS disease, multiple sclerosis, amyotrophic lateral sclerosis, Parkinson's disease and other neurological disorders, septic shock, sepsis, stroke, osteoporosis, osteoarthritis, ischemia reperfusion injury, cardiovascular disease, kidney disease, liver disease, ischemic injury, myocardial infarction, hypotension, hypertension, AIDS, myelodysplastic syndromes and other haematologic abnormalities, aplastic anaemia, male baldness pattern and bacterial, protozoal, fungal and viral infections related to SAP-4 polypeptide activity. Methods of identifying agonistic, antagonistic/inhibitors are also provided, as well as diagnostic assays for detecting diseases associated with inappropriate SAP-4 activity or levels.

Sequence 3099 BP: 769 A: 908 C: 813 G: 607 T: 0 U: 2 Other: XX

Query Match 100.0%; Score 1917; DB 2; Length 3099;

Query Match	100.0%;	Score	1317;
Best Local Similarity	100.0%;	Pred. No.	0;

Matches 1917; Conservative 0; Mismatches

THE UNIVERSITY OF CHICAGO

QY
1 ATGCTACTGCCACTGCTGCTGCTGCTGCTGCTG

[illegible]

51 ATGCTACTGCCACTGCTGCTGTCTCGCTGCTG
Db

61 TTCTTGGATACGAGTGGCAGGAGTCAGTGAATGCTG

U9
91 TCTGGATACGAGTGCAGGAGTCAGTGGATGGTGG

Db 111 TTCTGGATACGAGTCAGGAGTCAGTCAGTCATGGTG

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